## Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

## Listing of Claims:

- 1. (Original) A bioadhesive pharmaceutical formulation comprising an active agent and a mucoadhesive carrier for the active agent, wherein the mucoadhesive carrier comprises a β-limit dextrin.
  - 2. (Canceled).
- 3. (Previously presented) A bioadhesive pharmaceutical formulation as claimed in Claim 1 in which the active agent is a pharmaceutically active agent.
- 4. (Currently amended) A bioadhesive pharmaceutical formulation as claimed in Claim 1 which is a buccal-melt type product.
- 5. (Original) A bioadhesive pharmaceutical formulation as claimed in Claim 4 which is a wafer.
- 6. (Previously presented) A bioadhesive pharmaceutical formulation as claimed in Claim 1 which is a powder for use in aerosol delivery formulations.
- 7. (Previously presented) A bioadhesive pharmaceutical formulation as claimed in Claim 1 which is a thin film.
- 8. (Previously presented) A bioadhesive pharmaceutical formulation as claimed in Claim 1 further including at least one carbohydrate.

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- 9. (Original) A bioadhesive pharmaceutical formulation as claimed in Claim 8 in which the at least one carbohydrate is a polysaccharide.
- 10. (Previously presented) A bioadhesive pharmaceutical formulation as claimed in Claim 8 in which the at least one carbohydrate is selected from the group consisting of alginate; pectin; and derivatives of alginate and pectin.
- 11. (Original) A bioadhesive pharmaceutical formulation as claimed in Claim 10 in which the alginate comprises between 1 and 50% of the formulation (w/w).
- 12. (Original) A bioadhesive pharmaceutical formulation as claimed in Claim 11 in which the alginate comprises between 10 and 30% of the formulation (w/w).
  - 13. (Canceled)
  - 14. (Canceled)
  - 15. (Canceled)
- 16. (Previously presented) A bioadhesive pharmaceutical formulation as claimed in Claim 1 in a form selected from the group consisting of particulate; capsule; tablet; freeze dried matrix; wafer; liquid; and thin film.
- 17. (Original) A nutritional product comprising β-limit dextrin in which the β-limit dextrin is a main energy source in the product.
  - 18. (Original) A nutritional product as claimed in Claim 17 which is an energy drink.
  - 19. (Previously presented) A nutritional product as claimed in Claim 17 which is a

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confectionery product.

- 20. (Canceled)
- 21. (Canceled)
- 22. (Canceled)
- 23. (Canceled).
- 24. (Canceled)
- 25. (Currently amended) A bioadhesive pharmaceutical formulation <u>as</u> claimed in Claim 1 23 in which the starch is a waxy starch.
- 26. (Previously presented) A method for delivering an active agent to a mucosal membrane of a mammal comprising administering to said mammal a bioadhesive formulation comprising said active agent and a mucoadhesive carrier for the active agent, wherein the mucoadhesive carrier comprises a β-limit dextrin.
- 27. (Previously presented) A method according to Claim 26, wherein the active agent is a breath freshener.
- 28. (Previously presented) A method according to Claim 27, wherein the formulation is a thin-film breath freshener.
- 29. (Previously presented) A method according to Claim 26, wherein the active agent is a pharmaceutically active agent.

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- 30. (Previously presented) A method of providing nutrition to a subject comprising administering to the subject a nutritional product comprising a β-limit dextrin as an energy source.
- 31. (Previously presented) A method according to Claim 30, wherein the \(\beta\)-limit dextrin is the main energy source in the product
- 32. (Previously presented) A method according to Claim 30, wherein the β-limit dextrin is a slow release energy source.
- 33. (Previously presented) A method according to Claim 30, wherein the nutritional product is an energy drink.
- 34. (Previously presented) A method according to Claim 30, wherein the β-iimit dextrin is obtainable by hydrolysing starch.
- 35. (Previously presented) A method according to Claim 34, wherein the β-limit dextrin is obtainable by hydrolysing starch with β-amylase.
- 36. (Previously presented) A bioadhesive pharmaceutical formulation according to claim 1, wherein said formulation is a lyophilized formulation.
- 37. (Previously presented) A method according to claim 26, wherein said formulation is a lyophilized formulation.

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